Effect of age on contrast sensitivity function: uniconular and binocular findings

J E ROSS,1 D D CLARKE,2 AND A J BRON1

From the 1Nuffield Laboratory of Ophthalmology, University of Oxford, and the
2Department of Experimental Psychology, University of Oxford, Oxford

SUMMARY Monocular and binocular contrast sensitivity function for a range of spatial frequencies was measured in two groups of subjects with normal vision. Statistically significant differences in performance between the younger group (age 20–30 years) and the older group (age 50–87 years) were found at all spatial frequencies sampled between 0.40 and 19.25 cpd. In the age range 50–87 years there was a linear decline in contrast sensitivity with age for medium and high spatial frequencies, but sensitivity for low spatial frequencies was independent of age.

In recent years a number of investigators have used contrast sensitivity function (CSF) as a supplementary measure of visual performance both in normal subjects1 2 and in patients.3–5 Contrast thresholds are measured for a number of different spatial frequencies (the number of repetitions of a grating pattern, usually sinusoidal, within a unit of distance), and the results are represented graphically by plotting the reciprocal of the contrast threshold (contrast sensitivity) as a function of spatial frequency. In this way the performance of the eye can be specified over a wide working range and gives a broader description of visual function than the more conventional tests of visual acuity. Measurement of contrast sensitivity has revealed visual loss in patients with cerebral lesions,6 multiple sclerosis,7 macular disease,8 glaucoma,9 10 cataract,10 and amblyopia.11

A number of the above disorders are prevalent among older patients in whom age-related changes in the eye are likely to influence the results of visual function tests. It is thought that age-related changes, such as those found with visual acuity measurement, are attributable not only to lenticular yellowing and senile miosis but also to neuronal and receptor loss in the visual pathway.12 The implication that the decline with age in human visual performance is the result of loss of neuronal quality and quantity receives support from other sources. Age-related changes in the morphology of human cones13 and a decrease in the neuron population density of the macular projection areas in the visual cortex14 have been reported. In a tachistoscopic study Walsh15 has shown very clearly that there is a decrease in the efficiency of central processing of visual images with age.

Recently age dependent losses of contrast sensitivity have been reported by investigators who have used gratings displayed on an oscilloscope9 16–20 and the Arden printed test.9 21 22 Unfortunately the results of these studies do not all agree. Some investigators report a loss of contrast sensitivity for all spatial frequencies,9 19 21 while others found losses for high and medium spatial frequencies only.16 17 20 22 One group found losses only at low and medium spatial frequencies.18 It is likely that this lack of agreement between studies is attributable to a number of procedural differences, to the type of equipment used, and to the small sample sizes in the majority of studies.

Four of the studies investigated visual performance in patients of more than 70 years.9 18–20 Undoubtedly the study of older normal subjects poses a number of difficulties for the investigator and a definition of ‘normal’ becomes more difficult with increasing age. However, the incidence rates for disorders such as cataract, glaucoma, and macular disease in which abnormal CSF has been reported increase after the age of 65 years,23 so that this is an age group for which it is particularly important to obtain normative data in order to be able to evaluate visual dysfunction in relation to age. We have measured CSF in 70 subjects with normal eyes to obtain age norms for clinical use and to form a database for studies of various eye
disorders. Fifty three of these subjects were aged 50 or more. The remaining 17 subjects were young adults who were included for comparison with the older group.

Materials and methods

Subjects. Seventy Caucasians aged between 20 and 87 years volunteered to take part in the study. None of the subjects had any previous experience in contrast sensitivity measurement techniques. The responses for two different age groups were examined. In the younger age group there were 17 subjects: 7 males, mean age 23 years (range 22–26 years), and 10 females, mean age 25 years (range 20–30 years). In the older group there were 53 subjects: 24 males, mean age 71 years (range 54–81 years), and 29 females, mean age 73 years (range 50–87 years).

Subjects were included in the study if: (a) There was no current eye disorder on clinical examination. (b) There was no previous history of eye disorder that might affect visual function, for example, amblyopia. (c) There was no family history of glaucoma or diabetes. (d) There was visual acuity of 6/9 (decimal 0.66) or better in each eye separately. (e) Optical correction was between –6 dioptres and +6 dioptres inclusive.

Cataract was excluded on the basis of the presence of a black lens opacity silhouetted against the red reflex on ophthalmoscopic examination. Any eye in which such an opacity was observed was excluded from the study. Severity of nuclear sclerosis was assessed in the slit beam on a scale of one to three. A score of one indicated the least sclerosis, and a score of three indicated the most sclerosis. Subjects with a score of two or less in each eye separately were included in the sample.

Glaucoma was excluded on the basis of a careful family history, examination of the optic disc, and applanation tonometry for those aged 70 or more. Subjects with intraocular pressure of less than 21 mmHg were considered to be non-glaucomatous. Senile macular degeneration was excluded on the basis of a morphological absence of macular degenerative change. All but seven of the subjects had visual acuity of 6/6 (1.00) or better in each eye. These seven were subjects in the older age group: five had visual acuity of 6/9 (0.66) in one eye and 6/6 (1.00) or better in the fellow eye, and two subjects aged 81 and 87 had visual acuity of 6/12 (0.50) in one and 6/9 (0.66) in the other. The two eyes with visual acuity of 6/12 (0.50) were excluded from the study. The apparent pupil size for all eyes was greater than 2.5 mm.

Apparatus. Distance visual acuity was measured at 6 m by means of an internally illuminated Snellen chart that conformed to the British standard 4274. The luminance of the chart was 125 cd/m². A forced choice method was used to obtain visual acuity measurements.

Stationary vertical sine wave gratings of variable spatial frequency were generated on a display oscilloscope by a two channel computer addressed microprocessor waveform generator. The contrast of the grating pattern was adjusted by a computer linked attenuator. Contrast is defined as $\frac{(L_{\text{max}}-L_{\text{min}})}{(L_{\text{max}}+L_{\text{min}})}$ where $L_{\text{max}}$ and $L_{\text{min}}$ are the maximum and minimum luminances respectively of the grating bars. A device wired in parallel with the microprocessor permitted the contrast of sine wave gratings to be faded on to the oscilloscope screen over a period of 0·5 s before the preprogrammed contrast level was achieved. This device overcomes the problem of stimulating transient mechanisms in the visual system at stimulus onset.

The stimulus area on the oscilloscope screen was rectangular, 30 cm x 20 cm, with a matt grey surround extending out to form an area 35 cm x 42 cm. The mean screen luminance was 300 cd/m² and was independent of the contrast of the spatial frequency being displayed. The frame rate was 100 Hz. Contrast values were linear for the six spatial frequencies which were used (0.4, 0.95, 2.88, 6.73, 12.70, 19.23 cycles per degree of visual angle (cpd)). Changes in contrast of the sine wave gratings and the spatial frequency were either made interactively with the computer or preprogrammed. The subject viewed the screen from a distance of 280 cm and rested his chin on an adjustable rest. The test was described, and the subject was asked to respond to the presence of a vertical grating pattern, however faint, by pressing a hand held buzzer.

A preliminary routine was employed with the use of the interactive mode to establish the approximate range of threshold values for the six spatial frequencies. The gratings were presented on the screen in a pseudo random order at various contrast levels. This sequence provided an opportunity for the subject to become familiar with the grating patterns. An estimation of the contrast range in which the subject’s threshold was likely to fall for each spatial frequency was obtained from these measurements and recorded. For each spatial frequency two levels of contrast were used to begin a double staircase sequence. One contrast value was above the predicted contrast threshold, and one contrast value was below.

After the preliminary trial had been completed two preprogrammed sequences were used: a double staircase technique, and the ‘up-down transformed response rule’ (UDTR) technique. The contrast level at which each double staircase was begun was
Effect of age on contrast sensitivity function: uniocular and binocular findings

Fig. 1 A series of responses for the UDTR rule. The change in contrast level is represented on the vertical axis. The trial number is shown on the horizontal axis, and three reversals R are shown. The average of the values of the contrast at three reversals gives the value of contrast threshold. ■ = positive response. □ = negative response.

determined by the response recorded in the preliminary trial. The choice of staircase represented on a given trial was made randomly by the computer. For the double staircase the contrast was increased after a negative response and decreased after a positive response. The contrast increment or decrement step was calculated for each presentation as a fraction (0.079) of the previous contrast values.

When the crossover point on the double staircase had been reached a single sequence of stimuli for each spatial frequency value was then presented to the subject (the UDTR rule). The sequence began at the level of the last recorded values for each of the two staircases corresponding to a given spatial frequency. As with the double staircase the contrast was varied in steps of a predetermined size (0.039), but no change in contrast level was made until one of the following patterns of results was obtained at any level:

Decrease contrast after: YYY or OYY
Increase contrast after: YYO, OYO, OOO, or YOO
Y=positive response
O=negative response

This set of responses converges on the stimulus level that gives approximately 70% of positive responses (Fig. 1). The threshold was computed from the mean of three reversals.

CSF measurements were made for the right, left, and both eyes of each subject. For the monocular viewing condition the non-viewing eye was covered with an eye patch. In order to assess the reliability of measurements 20 of the subjects (10 males and 10 females) were retested approximately 24 hours after the initial test session.

Results

Monocular and binocular CSF results for the two groups were characterised by a smooth curve which peaked between 2 and 5 cpd and was typically attenuated at low and high spatial frequencies (Fig. 2). Monocular sensitivity was consistently lower than binocular sensitivity for all spatial frequencies. No statistically significant sex differences were found, and so data for male and female subjects were analysed together. In order to assess the reliability of responses the correlation coefficients derived from CSF values on a test-retest basis were examined. For each spatial frequency the correlation between the two test sessions exceeded 0.80 (p<0.001).

A one way analysis of variance was performed to determine whether the age related contrast sensitivity differences between the younger and older groups of observers were statistically significant (Fig. 2). For both viewing conditions there were significant
Table 1  Contrast sensitivity function: one way analysis of variance between young and old observers for monocular and binocular viewing

<table>
<thead>
<tr>
<th>Spatial frequency (cpd)</th>
<th>Group (age 50–87)</th>
<th>Subjects (age 20–30)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Group</td>
<td>Coefficient of variation (SD/mean × 100)</td>
</tr>
<tr>
<td></td>
<td>mean value</td>
<td>SD</td>
</tr>
<tr>
<td>Monocular viewing</td>
<td></td>
<td></td>
</tr>
<tr>
<td>0-40</td>
<td>1-435</td>
<td>0-187</td>
</tr>
<tr>
<td>0-95</td>
<td>2-051</td>
<td>0-173</td>
</tr>
<tr>
<td>2-88</td>
<td>2-542</td>
<td>0-215</td>
</tr>
<tr>
<td>6-73</td>
<td>2-284</td>
<td>0-312</td>
</tr>
<tr>
<td>12-70</td>
<td>1-736</td>
<td>0-380</td>
</tr>
<tr>
<td>19-25</td>
<td>1-252</td>
<td>0-284</td>
</tr>
<tr>
<td>Binocular viewing</td>
<td></td>
<td></td>
</tr>
<tr>
<td>0-40</td>
<td>1-494</td>
<td>0-212</td>
</tr>
<tr>
<td>0-95</td>
<td>2-098</td>
<td>0-171</td>
</tr>
<tr>
<td>2-88</td>
<td>2-630</td>
<td>0-222</td>
</tr>
<tr>
<td>6-73</td>
<td>2-382</td>
<td>0-291</td>
</tr>
<tr>
<td>12-70</td>
<td>1-836</td>
<td>0-344</td>
</tr>
<tr>
<td>19-25</td>
<td>1-350</td>
<td>0-310</td>
</tr>
</tbody>
</table>

Table 2  Regression analysis of monocular contrast sensitivity on age

<table>
<thead>
<tr>
<th>CPD</th>
<th>Slope</th>
<th>Intercept</th>
<th>r</th>
<th>p</th>
</tr>
</thead>
<tbody>
<tr>
<td>0-40</td>
<td>0.00307</td>
<td>1.0</td>
<td>&gt;0.05</td>
<td></td>
</tr>
<tr>
<td>0-95</td>
<td>-0.0285</td>
<td>2.10</td>
<td>&lt;0.05</td>
<td>&lt;0.05</td>
</tr>
<tr>
<td>2-88</td>
<td>-0.01203</td>
<td>2.80</td>
<td>&lt;0.001</td>
<td>&lt;0.05</td>
</tr>
<tr>
<td>6-73</td>
<td>-0.01472</td>
<td>2.66</td>
<td>&lt;0.001</td>
<td>&lt;0.05</td>
</tr>
<tr>
<td>12-70</td>
<td>-0.02017</td>
<td>2.20</td>
<td>&lt;0.001</td>
<td>&lt;0.05</td>
</tr>
<tr>
<td>19-25</td>
<td>-0.01578</td>
<td>1.61</td>
<td>&lt;0.01</td>
<td></td>
</tr>
</tbody>
</table>

The responses of the older age group were examined in more detail. A linear regression analysis of contrast sensitivity on age was made for all spatial frequencies. There was a significant decrease in contrast sensitivity with age for all but the two lowest spatial frequencies. The amount of scatter around the regression line was similar for monocular and binocular viewing, the least amount of scatter being found for a spatial frequency of 2-88 cpd in both cases (Table 2).

The older patients were divided into five age groups in order to examine the change with age in visual acuity measurement. However, entry to the study required visual acuity in each eye of 6/9 or better, and so the range of acuities sampled was necessarily limited. The mean visual acuity values for each age group are given in Table 3. There was a statistically significant difference among the means of the five age groups. The main effect for age was significant (F(4,99)=16-40, p<0-01). The relationship between a given level of visual acuity (6/5, 1-2, 6/6 0.80, or 6/7-5 or 6/6) and contrast sensitivity function was examined. A one way analysis of variance was performed, and no significant difference was found among the mean CSF values for the four acuity levels. However, for the highest two spatial frequencies there was a rank order fall of contrast sensitivity with each level of visual acuity.

Table 3  Mean visual acuity values for older subjects

<table>
<thead>
<tr>
<th>Age group (years)</th>
<th>Group mean: log minimum angle of resolution</th>
<th>Snellen equivalent</th>
</tr>
</thead>
<tbody>
<tr>
<td>48–60 n=20 eyes</td>
<td>1.9</td>
<td>6/5</td>
</tr>
<tr>
<td>61–70 n=22 eyes</td>
<td>0.0</td>
<td>6/6</td>
</tr>
<tr>
<td>71–75 n=18 eyes</td>
<td>0.0073</td>
<td>6/6</td>
</tr>
<tr>
<td>76–80 n=24 eyes</td>
<td>0.04</td>
<td>&gt;6/7-5</td>
</tr>
<tr>
<td>81 and over n=20 eyes</td>
<td>0.10</td>
<td>6/7-5</td>
</tr>
</tbody>
</table>
Discussion

We measured contrast sensitivity function in two
groups of normal subjects, first, to make a com-
parison of results between the older and the younger
subjects, and second, to examine the pattern of
results over a broad age range in 53 older subjects.
The study was conducted to obtain age norms for
clinical use and to form a database for studies of
various eye disorders. In addition to monocular test-
ing binocular assessments were performed in order to
relate the results of visual function tests to visual
performance as part of a wider study.29

The results showed that older observers had
reduced contrast sensitivity for all spatial frequencies
when compared with their younger counterparts. This
was particularly marked for medium and high spatial
frequencies. In the age range 50 to 87 years there was
a linear decline in CSF with age for medium and high
spatial frequencies. Within this age range the
responses to low spatial frequencies appeared to be
independent of age. These findings are in close agree-
ment with many of the existing data obtained using
oscilloscope generated gratings16 17 and printed
tests,19 22 but are opposed to those of Sekuler and
Hutman,18 who found low but not high spatial fre-
quency changes with increasing age.

Vaegan and Halliday1 consider that the most
important reason for Sekuler and Hutman’s18 con-
trary findings is that their elderly sample was not
representative, because 10 subjects had 6/6 (1·00)
vision or better in the tested eye. They argue that the
subjects came from a group with particularly high
visual acuity for the age range who do not have the
same visual performance for lower spatial fre-
quencies. However, the older subjects used in the
present study by reason of selection criteria suffer
from the same bias, in that the observed Snellen
visual acuities are better than one might expect in a
representative sample of people in the older age
range. Yet no age dependent changes for low spatial
frequencies were observed. In a more recent study by
the same group using a larger sample29 the authors
found that sensitivity to medium and high spatial
frequencies decreased with age after about 40 years.
As the same method to determine contrast thresholds
was used in both studies, it is possible that the age
dependent low frequency attenuation of sensitivity in
the first study18 is attributable to biases introduced by
using a small population sample.

Thus there appears to be little evidence for
preferential sensitivity losses at low spatial fre-
quencies with aging. It is well known that degra-
dation of any mechanical system, such as radios or hi-fi
sets, results primarily in the loss of high frequency
information. The visual system reacts in a similar
manner to degraded images.30 It is the low spatial
frequency information which conveys the major in-
formation about everyday objects as has been
demonstrated in studies of face recognition31 32 and
letter recognition.33 So while it seems likely that aging
changes at cellular level occur for neurones mediating
both high and low spatial frequency information,
these changes are less likely to show themselves in
response to low frequency stimuli.

The question whether senile miosis contributes to
the reduction of CSF with age has been raised by a
number of investigators.19 20 Derefeldt, Lennerstrand,
and Lundh17 do not give pupil sizes in their publica-
tion, but in the studies of Sekuler and Hutman16 and
McGrath and Morrison19 the pupil sizes were greater
than 2 mm as were those in the present study. Senile
miosis may not affect contrast sensitivity as such.
Other contributory factors may be involved. How-
ever, Woodhouse34 has shown that in the presence of
clear optical media, and with screen luminance
values above 34 cd/m², the difference between spatial
frequencies resolved at pupil diameters between 2
and 4 mm is minimal at both high (0·97) and low
(0·11) contrasts. In the present study the subjects had
clear media, the screen luminance was well above 34
cd/m², and the minimum pupil size was greater than
2·5 mm. The limiting effects of diffraction can be
dismissed at this pupil size, which suggests that in this
study senile miosis is unlikely to be a contributory
factor in the reduction of contrast sensitivity with
age.

It appears from the present data that, in common
with a number of other visual functions, CSF declines
with age and in particular at the medium and high
spatial frequency end of the curve. The existing data
are commensurate with this finding, though lack of
standardisation of stimuli and psychophysical pro-
cedures from laboratory to laboratory has resulted in
variations as to the magnitude and exact location of
the sensitivity loss on the contrast sensitivity curve.
Responses to stimuli at the low spatial frequency end
of the spectrum were independent of age in the
subjects over 50 years old in the present study and in
the majority of other studies. When pathological
status is assessed there are obvious advantages for the
use of stimuli which are relatively immune to aging
effects in the visual system. An added advantage of
the use of low spatial frequencies is that the effects of
optical aberration are minimal at this end of the CSF
curve.1

There is great potential for the application of CSF
testing in a number of visual disorders. In the present
study a series of age norms is presented for six spatial
frequencies with the use of a rigorous testing scheme.
Though it is clear that different investigators will
present findings based on the particular arrange-
ments available to them, there is a need to standardise approaches in order to make CSF testing more generally useful and available.

This research was supported by grant No 81/2 from Oxford Regional Health Authority.

References


J E Ross, D D Clarke, and A J Bron

Effect of age on contrast sensitivity function: uniocular and binocular findings.

J E Ross, D D Clarke and A J Bron

doi: 10.1136/bjo.69.1.51

Updated information and services can be found at:
http://bjo.bmj.com/content/69/1/51

These include:

Email alerting service

Receive free email alerts when new articles cite this article. Sign up in the box at the top right corner of the online article.

Notes

To request permissions go to:
http://group.bmj.com/group/rights-licensing/permissions

To order reprints go to:
http://journals.bmj.com/cgi/reprintform

To subscribe to BMJ go to:
http://group.bmj.com/subscribe/